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INK JET RECORDING ELEMENT

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INK JET RECORDING ELEMENT

CROSS REFERENCE TO RELATED APPLICATION

Reference is made to commonly assigned, co-pending U.S. Patent Application Serial Number: 10/008777 by Merkel et al., filed on ~~3~~ December 4, 2001 ~~of even date~~ herewith (Docket 83746), entitled "Ink Jet Printing Method".

FIELD OF THE INVENTION

This invention relates to an ink jet recording element. More particularly, this invention relates to a porous ink jet recording element containing porous polymeric particles.

BACKGROUND OF THE INVENTION

In a typical ink jet recording or printing system, ink droplets are ejected from a nozzle at high speed towards a recording element or medium to produce an image on the medium. The ink droplets, or recording liquid, generally comprise a recording agent, such as a dye or pigment, and a large amount of solvent. The solvent, or carrier liquid, typically is made up of water, an organic material such as a monohydric alcohol, a polyhydric alcohol or mixtures thereof.

An ink jet recording element typically comprises a support having on at least one surface thereof an ink-receiving or image-forming layer, and includes those intended for reflection viewing, which have an opaque support, and those intended for viewing by transmitted light, which have a transparent support.

While a wide variety of different types of image-recording elements for use with ink jet devices have been proposed heretofore, there are many unsolved problems in the art and many deficiencies in the known products which have limited their commercial usefulness.

It is well known that in order to achieve and maintain photographic-quality images on such an image-recording element, an ink jet recording element must:

- Be readily wetted so there is no puddling, i.e., coalescence of adjacent ink dots, which leads to non-uniform density

- Exhibit no image bleeding
- Absorb high concentrations of ink and dry quickly to avoid elements blocking together when stacked against subsequent prints or other surfaces
- Exhibit no discontinuities or defects due to interactions between the support and/or layer(s), such as cracking, repellencies, comb lines and the like
- Not allow unabsorbed dyes to aggregate at the free surface causing dye crystallization, which results in bloom or bronzing effects in the imaged areas
- Have an optimized image fastness to avoid fade from contact with water or radiation by daylight, tungsten light, or fluorescent light

An ink jet recording element that simultaneously provides an almost instantaneous ink dry time and good image quality is desirable. However, given the wide range of ink compositions and ink volumes that a recording element needs to accommodate, these requirements of ink jet recording media are difficult to achieve simultaneously.

Ink jet recording elements are known that employ porous or non-porous single layer or multilayer coatings that act as suitable image-receiving layers on one or both sides of a porous or non-porous support. Recording elements that use non-porous coatings typically have good image quality but exhibit poor ink dry time. Recording elements that use porous coatings exhibit superior dry times, but typically have poorer image quality and are prone to cracking.

A problem with known ink jet recording elements that employ porous single layer or multilayer coatings that act as suitable image-receiving layer is dye stability during storage. In particular, dyes printed on to an ink jet receiver element tend to fade due to exposure to ozone which is present in the atmosphere.

Another problem with ink jet recording elements that employ porous single layer or multilayer coatings that act as suitable image-receiving layers is image stability under high humidity storage conditions. In particular, dyes tend to migrate through the image receiving layer during storage since the

dye image receiving layer is hydrophilic and tends to absorb water from the atmosphere.

Copending U.S. Patent Application S.N. 09/608,466, filed June 30, 2000, relates to an jet recording element wherein the image-receiving layer contains porous polymeric particles in a polymeric binder. However, there is a problem with this element in that during preparation of the coating solution, agglomeration of the polymeric particles occurs, which when coated, results in an element having low gloss.

JP 09207430, JP 08324101 and JP 2000/239578 relate to porous image-receiving layers for ink jet recording elements containing inorganic particles and a poly(vinyl alcohol) having various degrees of hydrolysis. However, there is a problem with these elements in that the references do not disclose the degree of hydrolysis for the poly(vinyl alcohol) necessary to provide good gloss and low cracking.

It is an object of this invention to provide an ink jet recording element that has a fast ink dry time. It is another object of this invention to provide an ink jet recording element that has good stability when exposed to ozone and high humidity conditions. It is another object of the invention to provide an ink jet recording element that has high gloss with minimal cracking.

SUMMARY OF THE INVENTION

These and other objects are achieved in accordance with the invention which comprises an ink jet recording element comprising a support having thereon an image-receiving layer comprising porous polymeric particles in a polymeric binder, the polymeric binder comprising poly(vinyl alcohol) having a degree of hydrolysis of at least about 95% and having a number average molecular weight of at least about 45,000.

By use of the invention, an ink jet recording element is obtained which has a good dry time and good stability when exposed to ozone and high humidity conditions, and has high gloss with minimal cracking.

DETAILED DESCRIPTION OF THE INVENTION

The support used in the ink jet recording element of the invention may be opaque, translucent, or transparent. There may be used, for example, plain papers, resin-coated papers, various plastics including a polyester resin such as poly(ethylene terephthalate), poly(ethylene naphthalate) and poly(ester diacetate), a polycarbonate resin, a fluorine resin such as poly(tetra-fluoro ethylene), metal foil, various glass materials, and the like. In a preferred embodiment, the support is paper or a voided plastic material. The thickness of the support employed in the invention can be from about 12 to about 500 μm , preferably from about 75 to about 300 μm .

The porous polymeric particles which are used in the invention are in the form of porous beads, porous irregularly shaped particles, or are aggregates of emulsion particles.

Suitable porous polymeric particles used in the invention comprise, for example, acrylic resins, styrenic resins, or cellulose derivatives, such as cellulose acetate, cellulose acetate butyrate, cellulose propionate, cellulose acetate propionate, and ethyl cellulose; polyvinyl resins such as polyvinyl chloride, copolymers of vinyl chloride and vinyl acetate and polyvinyl butyral, polyvinyl acetal, ethylene-vinyl acetate copolymers, ethylene-vinyl alcohol copolymers, and ethylene-allyl copolymers such as ethylene-allyl alcohol copolymers, ethylene-allyl acetone copolymers, ethylene-allyl benzene copolymers, ethylene-allyl ether copolymers, ethylene acrylic copolymers and polyoxy-methylene; polycondensation polymers, such as, polyesters, including polyethylene terephthalate, polybutylene terephthalate, polyurethanes and polycarbonates.

In a preferred embodiment of the invention, the porous polymeric particles are made from a styrenic or an acrylic monomer. Any suitable ethylenically unsaturated monomer or mixture of monomers may be used in making such styrenic or acrylic polymer. There may be used, for example, styrenic compounds, such as styrene, vinyl toluene, p-chlorostyrene, vinylbenzylchloride or vinyl naphthalene; or acrylic compounds, such as methyl acrylate, ethyl acrylate, n-butyl acrylate, n-octyl acrylate, 2-chloroethyl acrylate, phenyl acrylate, methyl- α -chloroacrylate, methyl methacrylate, ethyl

methacrylate, butyl methacrylate; and mixtures thereof. In another preferred embodiment, methyl methacrylate or ethylene glycol dimethacrylate is used.

In a preferred embodiment of the invention, the porous polymeric particles are crosslinked. They may have a degree of crosslinking of about 27 mole % or greater, preferably about 50 mole %, and most preferably about 100 mole %. The degree of crosslinking is determined by the mole % of multifunctional crosslinking monomer which is incorporated into the porous polymeric particles.

Typical crosslinking monomers which may be used in making the porous polymeric particles employed in the invention are aromatic divinyl compounds such as divinylbenzene, divinyl naphthalene or derivatives thereof; diethylene carboxylate esters and amides such as ethylene glycol dimethacrylate, diethylene glycol diacrylate, and other divinyl compounds such as divinyl sulfide or divinyl sulfone compounds. Divinylbenzene and ethylene glycol dimethacrylate are especially preferred.

The porous polymeric particles used in this invention can be prepared, for example, by pulverizing and classification of porous organic compounds, by emulsion, suspension, and dispersion polymerization of organic monomers, by spray drying of a solution containing organic compounds, or by a polymer suspension technique which consists of dissolving an organic material in a water immiscible solvent, dispersing the solution as fine liquid droplets in aqueous solution, and removing the solvent by evaporation or other suitable techniques. The bulk, emulsion, dispersion, and suspension polymerization procedures are well known to those skilled in the polymer art and are taught in such textbooks as G. Odian in "Principles of Polymerization", 2nd Ed. Wiley (1981), and W.P. Sorenson and T.W. Campbell in "Preparation Method of Polymer Chemistry", 2nd Ed, Wiley (1968).

Techniques to synthesize porous polymer particles are taught, for example, in U.S. Patents 5,840,293; 5,993,805; 5,403,870; and 5,599,889, and Japanese Kokai Hei 5[1993]-222108, the disclosures of which are hereby incorporated by reference. For example, an inert fluid or porogen may be mixed with the monomers used in making the porous polymer particles. After

polymerization is complete, the resulting polymeric particles are, at this point, substantially porous because the polymer has formed around the porogen thereby forming the pore network. This technique is described more fully in U.S. Patent 5,840,293 referred to above. Thus, the porosity of the porous polymeric particles is achieved by mixing a porogen with the monomers used to make the polymeric particles, dispersing the resultant mixture in water, and polymerizing the monomers to form the porous polymeric particles.

A preferred method of preparing the porous polymeric particles used in this invention includes forming a suspension or dispersion of ethylenically unsaturated monomer droplets containing the crosslinking monomer and a porogen in an aqueous medium, polymerizing the monomer to form solid, porous polymeric particles, and optionally removing the porogen by vacuum stripping. The particles thus prepared have a porosity as measured by a specific surface area of about 35 m²/g or greater, preferably 100 m²/g or greater. The surface area is usually measured by B.E.T. nitrogen analysis known to those skilled in the art.

The porous polymeric particles may be covered with a layer of colloidal inorganic particles as described in U.S. Patents 5,288,598; 5,378,577; 5,563,226 and 5,750,378, the disclosures of which are incorporated herein by reference. The porous polymeric particles may also be covered with a layer of colloidal polymer latex particles as described in U.S. Patent 5,279,934, the disclosure of which is incorporated herein by reference.

The porous polymeric particles used in this invention have a median diameter less than about 10 µm, preferably less than about 1 µm, and most preferably less than about 0.6 µm. Median diameter is defined as the statistical average of the measured particle size distribution on a volume basis. For further details concerning median diameter measurement, see T. Allen, "Particle Size Measurement", 4th Ed., Chapman and Hall, (1990).

As noted above, the polymeric particles used in the invention are porous. By porous is meant particles which either have voids or are permeable to liquids. These particles can have either a smooth or a rough surface.

The image-receiving layer of the ink jet recording element of the invention may contain a surfactant. Suitable surfactants include anionic surfactants or cationic surfactants.

As noted above, the poly(vinyl alcohol) employed in the invention
5 has a degree of hydrolysis of at least about 95% and has a number average molecular weight of at least about 45,000. In a preferred embodiment of the invention, the poly(vinyl alcohol) has a degree of hydrolysis of at least about 98%. In another preferred embodiment of the invention, the poly(vinyl alcohol) has a number average molecular weight of from about 70,000 to about 105,000.
10 Commercial embodiments of such a poly(vinyl alcohol) are Gohsenol ® AH-22, Gohsenol ® AH-26 and Gohsenol ® AH-17 from Nippon Gohsei.

The image-receiving layer may also contain additives such as pH-modifiers like nitric acid, cross-linkers, rheology modifiers, surfactants, UV-absorbers, biocides, lubricants, water-dispersible latexes, mordants, dyes, optical
15 brighteners etc.

The image-receiving layer may be applied to one or both substrate surfaces through conventional pre-metered or post-metered coating methods such as blade, air knife, rod, roll, slot die, curtain, slide, etc. The choice of coating process would be determined from the economics of the operation and in turn,
20 would determine the formulation specifications such as coating solids, coating viscosity, and coating speed.

The image-receiving layer thickness may range from about 5 to about 100 μm , preferably from about 10 to about 50 μm . The coating thickness required is determined through the need for the coating to act as a sump for
25 absorption of ink solvent. The image-receiving layer of this invention contains from about 0.20 to about 10.0 g/m^2 of polymeric binder, preferably from about 0.40 to about 5.0 g/m^2 , and about 1.5 to about 60 g/m^2 of porous polymeric particles, preferably from about 3.0 to about 30 g/m^2 .

Ink jet inks used to image the recording elements of the present
30 invention are well-known in the art. The ink compositions used in ink jet printing typically are liquid compositions comprising a solvent or carrier liquid, dyes or pigments, humectants, organic solvents, detergents, thickeners, preservatives, and

the like. The solvent or carrier liquid can be solely water or can be water mixed with other water-miscible solvents such as polyhydric alcohols. Inks in which organic materials such as polyhydric alcohols are the predominant carrier or solvent liquid may also be used. Particularly useful are mixed solvents of water and polyhydric alcohols. The dyes used in such compositions are typically water-soluble direct or acid type dyes. Such liquid compositions have been described extensively in the prior art including, for example, U.S. Patents 4,381,946; 4,239,543 and 4,781,758, the disclosures of which are hereby incorporated by reference.

Although the recording elements disclosed herein have been referred to primarily as being useful for ink jet printers, they also can be used as recording media for pen plotter assemblies. Pen plotters operate by writing directly on the surface of a recording medium using a pen consisting of a bundle of capillary tubes in contact with an ink reservoir.

The following example further illustrates the invention.

EXAMPLE

The following elements were prepared with the image-receiving layer as described:

Element 1 of the Invention

A 10% by weight solution of water, borax (sodium tetraborate decahydrate) and a sulfonated polyester dispersion AQ29® (Eastman Chemical Co.) with a coating surfactant Olin 10G®, with the borax to polyester binder ratio being 33:67, was rod coated on a corona-discharge treated resin coated paper for a total dry lay-down of 1.5 g/m², giving a dry lay-down of borax of 0.5 g/m² and a polyester binder dry lay-down of 1.0 g/m². The subbing layer coating was dried in a oven at 40° C for 20 minutes.

A second solution at about 18% by weight comprised of porous polymeric particles, poly(ethylene glycol dimethacrylate), and a poly(vinyl alcohol) binder, AH-26 from Nippon Gohsei, where the ratio of porous polymer particles to PVA was about 80:20, was blade coated over the subbing layer to a

dry lay-down of about 40 g/m² and dried at 40°C for about 20 minutes to provide an image-receiving layer.

The number average molecular weight of the poly(vinyl alcohol) listed in Table 2 was estimated from the viscosity of a 4% aqueous solution according to a table provided by a commercial manufacturer of poly(vinyl alcohol). The degree of hydrolysis of the poly(vinyl alcohol) was obtained from the manufacturer.

Element 2 of the Invention

10 This element was prepared the same as Element 1 except that the poly(vinyl alcohol) in the image-receiving layer was AH-22 from Nippon Gohsei.

Element 3 of the Invention

15 This element was prepared the same as Element 1 except that the poly(vinyl alcohol) in the image-receiving layer was AH-17 from Nippon Gohsei.

Control Element C-1 (Low m.w. PVA and low degree of hydrolysis)

20 This element was prepared the same as Element 1 except that the poly(vinyl alcohol) in the image-receiving layer was AL-06 from Nippon Gohsei.

Control Element C-2 (Low degree of hydrolysis)

This element was prepared the same as Element 1 except that the poly(vinyl alcohol) in the image-receiving layer was GH-23 from Nippon Gohsei.

25 Control Element C-3 (Low degree of hydrolysis)

This element was prepared the same as Element 1 except that the poly(vinyl alcohol) in the image-receiving layer was GH-17 from Nippon Gohsei.

Control Element C-4 (Low degree of hydrolysis)

30 This element was prepared the same as Element 1 except that the poly(vinyl alcohol) in the image-receiving layer was KH-20 from Nippon Gohsei.

Control Element C-5 (Low degree of hydrolysis)

This element was prepared the same as Element 1 except that the poly(vinyl alcohol) in the image-receiving layer was KH-17 from Nippon Gohsei.

5 Testing

Each element was imaged using an Epson 870 ink jet printer and ink jet inks, Cartridge No.T007 (black) and T008 (color), and then rated for cracking according to Table 1.

10

Table 1

<u>Rating</u>	<u>Cracking Observations</u>
1	No visible cracks under magnification
2	Slight micro-cracks under 10X magnification
3	Very slight visible cracks under no magnification
4	Heavy cracking, some flaking
5	Heavy cracking, coating flaking off

Each element was then measured for 60 degree gloss, using a Gardner Gloss meter. The average gloss of cyan, magenta, yellow, red, blue, green, black, and D-min was recorded in Table 2. Average gloss level of greater

15 than about 35 is acceptable.

Table 2

<u>Element</u>	<u>PVA</u>	<u>Degree of Hydrolysis</u>	<u>Approximate Number Average m.w.</u>	<u>Cracking Rating</u>	<u>Gloss</u>
1	AH-26	98	90,000-100,000	2	45
2	AH-22	98	80,000-90,000	2	43
3	AH-17	98	60,000-65,000	4	38
C-1	AL-06	92	25,000-30,000	4	10
C-2	GH-23	88	80,000-90,000	2	10
C-3	GH-17	88	60,000-65,000	3	12
C-4	KH-20	80	70,000-80,000	3	15
C-5	KH-17	80	65,000-70,000	4	15

The above results show that the elements according to the invention having a poly(vinyl alcohol) with the degree of hydrolysis of at least about 95% and a number average molecular weight of at least about 45,000 all provide good gloss as compared to the control elements. In addition, the elements
5 according to invention with a degree of hydrolysis of at least about 95% and an number average molecular weight of at least about 70,000 provide both in good gloss and low cracking as compared to the control elements.

This invention has been described with particular reference to preferred embodiments thereof but it will be understood that modifications can be
10 made within the spirit and scope of the invention.

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